COMMON TOXICITY CRITERIA (CTC)

		Gr	ade		
Toxicity	0	1	2	3	4
		ALLERGY/IM	IMUNOLOGY		
Allergic reaction/ hypersensitivity (including drug fever)	none	transient rash, drug fever < 38°C (<100.4°F)	urticaria, drug fever ≥ 38°C (≥100.4°F), and/or asymptomatic bronchospasm	symptomatic bronchospasm, requiring parenteral medication(s), with or without urticaria; allergy-related edema/angioedema	anaphylaxis
Note: Isolated urticaria, in the	absence of other r	nanifestations of an allergic or hy	persensitivity reaction, is gr	aded in the DERMATOLO	GY/SKIN category.
Allergic rhinitis (including sneezing, nasal stuffiness, postnasal drip)	none	mild, not requiring treatment	moderate, requiring treatment	-	-
Autoimmune reaction	none	serologic or other evidence of autoimmune reaction but patient is asymptomatic (e.g., vitiligo), all organ function is normal and no treatment is required	evidence of autoimmune reaction involving a non- essential organ or function (e.g., hypothyroidism), requiring treatment other than immunosuppressive drugs	reversible autoimmune reaction involving function of a major organ or other toxicity (e.g., transient colitis or anemia), requiring short-term immunosuppressive treatment	autoimmune reaction causing major grade 4 organ dysfunction; progressive and irreversible reaction; long-term administration of high- dose immuno- suppressive therapy required
Also consider Hypothyroidism		obin, Hemolysis.			
Serum sickness	none	-	-	present	<u>-</u>
Urticaria is graded in the DER reaction, grade as Allergic rea		IN category if it occurs as an isol vity above.	ated symptom. If it occurs v	with other manifestations of	allergic or hypersensitivity
Vasculitis	none	mild, not requiring treatment	symptomatic, requiring medication	requiring steroids	ischemic changes or requiring amputation
Allergy/Immunology-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		AUDITORY	Y/HEARING		
Conductive hearing loss is gra	ded as Middle ear	/hearing in the AUDITORY/HEA	ARING category.		
Earache is graded in the PAIN	category.				
External auditory canal	normal	external otitis with erythema or dry desquamation	external otitis with moist desquamation	external otitis with discharge, mastoiditis	necrosis of the canal soft tissue or bone
Note: Changes associated with	n radiation to exter	rnal ear (pinnae) are graded under	Radiation dermatitis in the	DERMATOLOGY/SKIN ca	ategory.

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		Gr	ade		
Toxicity	0	1	2	3	4
Inner ear/hearing	normal	hearing loss on audiometry only	tinnitus or hearing loss, not requiring hearing aid or treatment	tinnitus or hearing loss, correctable with hearing aid or treatment	severe unilateral or bilateral hearing loss (deafness), not correctable
Middle ear/hearing	normal	serous otitis without subjective decrease in hearing	serous otitis or infection requiring medical intervention; subjective decrease in hearing; rupture of tympanic membrane with discharge	otitis with discharge, mastoiditis or conductive hearing loss	necrosis of the canal soft tissue or bone
Auditory/Hearing-Other (Specify,)	normal	mild	moderate	severe	life-threatening or disabling
		BLOOD/BON	E MARROW		
Bone marrow cellularity	normal for age	mildly hypocellular or 25% reduction from normal cellularity for age	moderately hypocellular or >25 - ≤ 50% reduction from normal cellularity for age or >2 but <4 weeks to recovery of normal bone marrow cellularity	severely hypocellular or >50 - ≤ 75% reduction in cellularity for age or 4 - 6 weeks to recovery of normal bone marrow cellularity	aplasia or >6 weeks to recovery of normal bone marrow cellularity
Normal ranges:					
children (≤ 18 years)	90% cellularity average				
younger adults (19-59)	60-70% cellularity average				
older adults (≥ 60 years)	50% cellularity average				
Note: Grade Bone marrow cell	ularity only for change	s related to treatment not dis	ease.		
CD4 count	WNL	< LLN - 500/mm ³	$200 - < 500/\text{mm}^3$	$50 - < 200/\text{mm}^3$	< 50/mm ³
Haptoglobin	normal	decreased	-	absent	-
Hemoglobin (Hgb)	WNL	< LLN - 10.0 g/dl < LLN - 100 g/L < LLN - 6.2 mmol/L	8.0 - < 10.0 g/dl 80 - < 100 g/L 4.9 - < 6.2 mmol/L	6.5 - < 8.0 g/dl 65 - 80 g/L 4.0 - < 4.9 mmol/L	< 6.5 g/dl < 65 g/L < 4.0 mmol/L
Note: The following criteria m	ay be used for leukemi	a studies or bone marrow int	ïltrative/myelophthisic proc	ess if the protocol so specifi	es.
For leukemia studies or bone marrow infiltrative/ myelophthisic processes	WNL	10 - <25% decrease from pretreatment	25 - <50% decrease from pretreatment	50 - <75% decrease from pretreatment	≥75% decrease from pretreatment
Hemolysis (e.g., immune hemolytic anemia, drug- related hemolysis, other)	none	only laboratory evidence of hemolysis [e.g., direct antiglobulin test (DAT, Coombs') schistocytes]	evidence of red cell destruction and ≥ 2gm decrease in hemoglobin, no transfusion	requiring transfusion and/or medical intervention (e.g., steroids)	catastrophic consequences of hemolysis (e.g., renal failure, hypotension, bronchospasm, emergency splenectomy)
Also consider Haptoglobin, Hg	gb.				

Toxicity Leukocytes (total WBC)	0 WNL	1	2	2	
Leukocytes (total WBC)	WNI.		-	3	4
	,,,,,	< LLN - 3.0 x 10 ⁹ /L < LLN - 3000/mm ³	$\geq 2.0 - < 3.0 \times 10^9 / L$ $\geq 2000 - < 3000 / mm^3$	$\geq 1.0 - < 2.0 \times 10^9 / L$ $\geq 1000 - < 2000 / mm^3$	< 1.0 x 10 ⁹ /L < 1000/mm ³
For BMT studies:	WNL	$\geq 2.0 - \langle 3.0 \times 10^9 / L$ $\geq 2000 - \langle 3000 / mm^3 \rangle$	$\geq 1.0 - \langle 2.0 \times 10^9 / L$ $\geq 1000 - \langle 2000 / mm^3 \rangle$	$\geq 0.5 - < 1.0 \times 10^9 / L$ $\geq 500 - < 1000 / mm^3$	<0.5 x 10 ⁹ /L <500/mm ³
Note: The following criteria usi	ing age, race and sex n	ormal values may be used fo	or pediatric studies if the pro	otocol so specifies.	
		³ 75 - <100% LLN	³ 50 - <75% LLN	³ 25 - 50% LLN	<25% LLN
Lymphopenia	WNL	<lln -="" 1.0="" 10<sup="" x="">9 /L <lln -="" 1000="" mm<sup="">3</lln></lln>	$\ge 0.5 - < 1.0 \times 10^9 / L$ $\ge 500 - < 1000 / mm^3$	<0.5 x 10 ⁹ /L <500/mm ³	-
Note: The following criteria usi	ing age, race, and sex r	normal values may be used f	or pediatric studies if the pr	otocol so specifies.	
		³ 75-<100%LLN	³ 50-<75%LLN	³ 25-<50%LLN	<25%LLN
Neutrophils/granulocytes (ANC/AGC)	WNL	$\geq 1.5 - \langle 2.0 \times 10^9 / L$ $\geq 1500 - \langle 2000 / mm^3 \rangle$	$\geq 1.0 - < 1.5 \times 10^9 / L$ $\geq 1000 - < 1500 / mm^3$	$\ge 0.5 - < 1.0 \times 10^9 / L$ $\ge 500 - < 1000 / mm^3$	< 0.5 x 10 ⁹ /L < 500/mm ³
For BMT:	WNL	$\geq 1.0 - < 1.5 \times 10^9 / L$ $\geq 1000 - < 1500 / mm^3$	$\ge 0.5 - < 1.0 \times 10^9 / L$ $\ge 500 - < 1000 / mm^3$	$\ge 0.1 - < 0.5 \times 10^9 / L$ $\ge 100 - < 500 / mm^3$	<0.1 x 10 ⁹ /L <100/mm ³
Note: The following criteria ma	ny be used for leukemia	studies or bone marrow inf	iltrative/myelophthisic proce	ess if the protocol so specific	es.
For leukemia studies or bone marrow infiltrative/ myelophthisic process	WNL	10 - <25% decrease from baseline	25 - <50% decrease from baseline	50 - <75% decrease from baseline	≥75% decrease from baseline
Platelets	WNL	< LLN - <75.0 x 10 ⁹ /L < LLN - 75000/mm ³	\geq 50.0 - < 75.0 x 10 ⁹ /L \geq 50000 - < 75000/mm ³	$\geq 10.0 - < 50.0 \times 10^9 / L$ $\geq 10000 - < 50000 / mm^3$	< 10.0 x 10 ⁹ /L < 10000/mm ³
For BMT:	WNL	≥50.0 - <75.0 x 10 ⁹ /L ≥50000 - <75000/mm ³	\geq 20.0 - <50.0 x 10 ⁹ /L \geq 20000 - <50000/mm ³	$\geq 10.0 - <20.0 \times 10^9 / L$ $\geq 10000 - <20000 / mm^3$	<10.0 x 10 ⁹ /L <10000/mm ³
Note: The following criteria ma	ny be used for leukemia	studies or bone marrow inf	iltrative/myelophthisic proce	ess if the protocol so specific	es.
For leukemia studies or bone marrow infiltrative/ myelophthisic process	WNL	10 - <25% decrease from baseline	25 - <50% decrease from baseline	50 - <75% decrease from baseline	≥75% decrease from baseline
Transfusion: Platelets	none	-	-	yes	platelet transfusions and other measures required to improve platelet increment; platelet transfusion refractoriness associated with life-threatening bleeding. (e.g., HLA or cross matched platelet transfusions)
For BMT: Also consider Platelets.	none	1 platelet transfusion in 24 hours	2 platelet transfusions in 24 hours	≥3 platelet transfusions in 24 hours	platelet transfusions and other measures required to improve platelet increment; platelet transfusion refractoriness associated with life-threatening bleeding. (e.g., HLA or cross matched platelet transfusions)

Grade							
Toxicity	0	1	2	3	4		
Transfusion: pRBCs	none	-	-	Yes	-		
For BMT:	none	≤2 u pRBC (≤15cc/kg) in 24 hours elective or planned	3 u pRBC (>15 $\leq 30cc/kg$) in 24 hours elective or planned	≥4 u pRBC (>30cc/kg) in 24 hours	hemorrhage or hemolysis associated with life-threatening anemia; medical intervention required to improve hemoglobin		
Also consider Hemoglobin.							
Blood/Bone Marrow-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling		
	CA	RDIOVASCULA	R (ARRHYTHM	IIA)			
Conduction abnormality/ Atrioventricular heart block	none	asymptomatic, not requiring treatment (e.g., Mobitz type I second-degree AV block, Wenckebach)	symptomatic, but not requiring treatment	symptomatic and requiring treatment (e.g., Mobitz type II second-degree AV block, third-degree AV block)	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)		
Nodal/junctional arrhythmia/dysrhythmia	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)		
Palpitations	none	present	-	-	-		
Note: Grade palpitations only i	n the absence of a doci	umented arrhythmia.					
Prolonged QTc interval (QTc > 0.48 seconds)	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)		
Sinus bradycardia	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)		
Sinus tachycardia	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment of underlying cause	-		
Supraventricular arrhythmias (SVT/atrial fibrillation/flutter)	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)		
Syncope (fainting) is graded in	the NEUROLOGY ca	tegory.					
Vasovagal episode	none	-	present without loss of consciousness	present with loss of consciousness	-		

	Grade							
Toxicity	0	1	2	3	4			
Ventricular arrhythmia (PVCs/bigeminy/trigeminy/ ventricular tachycardia)	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic and requiring treatment	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)			
Cardiovascular/ Arrhythmia-Other (Specify,)	none	asymptomatic, not requiring treatment	symptomatic, but not requiring treatment	symptomatic, and requiring treatment of underlying cause	life-threatening (e.g., arrhythmia associated with CHF, hypotension, syncope, shock)			
		CARDIOVASCU	LAR (GENERAL)				
Acute vascular leak syndrome	absent	-	symptomatic, but not requiring fluid support	respiratory compromise or requiring fluids	life-threatening; requiring pressor support and/or ventilatory support			
Cardiac- ischemia/infarction	none	non-specific T-wave flattening or changes	asymptomatic, ST- and T- wave changes suggesting ischemia	angina without evidence of infarction	acute myocardial infarction			
Cardiac left ventricular function	normal	asymptomatic decline of resting ejection fraction of ≥ 10% but < 20% of baseline value; shortening fraction ≥ 24% but < 30%	asymptomatic but resting ejection fraction below LLN for laboratory or decline of resting ejection fraction ≥ 20% of baseline value; < 24% shortening fraction	CHF responsive to treatment	severe or refractory CHF or requiring intubation			
CNS cerebrovascular ischemi	a is graded in the	NEUROLOGY category.						
Cardiac troponin I (cTnI)	normal	-	-	levels consistent with unstable angina as defined by the manufacturer	levels consistent with myocardial infarction as defined by the manufacturer			
Cardiac troponin T (cTnT)	normal	≥ 0.03 - < 0.05 ng/ml	≥ 0.05 - < 0.1 ng/ml	≥ 0.1 - < 0.2 ng/ml	≥ 0.2 ng/ml			
Edema	none	asymptomatic, not requiring therapy	symptomatic, requiring therapy	symptomatic edema limiting function and unresponsive to therapy or requiring drug discontinuation	anasarca (severe generalized edema)			
Hypertension	none	asymptomatic, transient increase by >20 mmHg (diastolic) or to > 150/100* if previously WNL; not requiring treatment	recurrent or persistent or symptomatic increase by > 20 mmHg (diastolic) or to > 150/100* if previously WNL; not requiring treatment	requiring therapy or more intensive therapy than previously	hypertensive crisis			

		Gr	ade		
Toxicity	0	1	2	3	4
Hypotension	none	changes, but not requiring therapy (including transient orthostatic hypotension)	requiring brief fluid replacement or other therapy but not hospitalization; no physiologic consequences	requiring therapy and sustained medical attention, but resolves without persisting physiologic consequences	shock (associated with acidemia and impairing vital organ function due to tissue hypoperfusion)
Also consider Syncope (fainti	ng).				
Note: Angina or MI is graded	d as Cardiac- ischemia	/infarction in the CARDIOVA	ASCULAR (GENERAL) ca	tegory.	
For pediatric patients, or three measurements		or less in infants up to 1 year	old and 70 mmHg or less in	n children older than 1 year	of age, use two successive
Myocarditis	none	-	-	CHF responsive to treatment	severe or refractory CHF
Operative injury of vein/artery	none	primary suture repair for injury, but not requiring transfusion	primary suture repair for injury, requiring transfusion	vascular occlusion requiring surgery or bypass for injury	myocardial infarction; resection of organ (e.g., bowel, limb)
Pericardial effusion/ pericarditis	none	asymptomatic effusion, not requiring treatment	pericarditis (rub, ECG changes, and/or chest pain)	physiologic consequences resulting from symptoms	tamponade (drainage or pericardial window required)
Peripheral arterial ischemia	none	-	brief episode of ischemia managed non- surgically and without permanent deficit	requiring surgical intervention	life-threatening or with permanent functional deficit (e.g., amputation)
Phlebitis (superficial)	none	-	present	-	-
Note: Injection site reaction i	is graded in the DERM	IATOLOGY/SKIN category.			
Thrombosis/embolism	is graded in the CARI	DIOVASCULAR (GENERAI	L) category.		
Syncope (fainting) is graded i	n the NEUROLOGY	category.			
Thrombosis/embolism	none	-	deep vein thrombosis, not requiring anticoagulant	deep vein thrombosis, requiring anticoagulant therapy	embolic event including pulmonary embolism
Vein/artery operative injury is	graded as Operative i	njury of vein/artery in the CA	ARDIOVASCULAR (GENE	ERAL) category.	
Visceral arterial ischemia (non-myocardial)	none	-	brief episode of ischemia managed non- surgically and without permanent deficit	requiring surgical intervention	life-threatening or with permanent functional deficit (e.g., resection of ileum)
Cardiovascular/ General-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling

Grade							
Toxicity	0	1	2	3	4		
		COAGU	LATION				
Note: See the HEMORRHAG	E category for grad	ling the severity of bleeding even	nts.				
DIC (disseminated intravascular coagulation)	absent	-	-	laboratory findings present with <u>no</u> bleeding	laboratory findings <u>and</u> bleeding		
Also grade Platelets.							
Note: Must have increased fibr	rin split products o	r D-dimer in order to grade as D	IC.				
Fibrinogen	WNL	≥0.75 - <1.0 x LLN	≥0.5 - <0.75 x LLN	≥0.25 - <0.5 x LLN	<0.25 x LLN		
Note: The following criteria m	ay be used for leuk	temia studies or bone marrow in	filtrative/myelophthisic proc	ess if the protocol so specifi	es.		
For leukemia studies:	WNL	<20% decrease from pretreatment value or LLN	≥20 - <40% decrease from pretreatment value or LLN	≥40 - <70% decrease from pretreatment value or LLN	<50 mg%		
Partial thromboplastin time (PTT)	WNL	> ULN - ≤ 1.5 x ULN	> 1.5 - ≤ 2 x ULN	>2 x ULN	-		
Phelbitis is graded in the CAR	DIOVASCULAR	(GENERAL) category.					
Prothrombin time (PT)	WNL	> ULN - ≤ 1.5 x ULN	> 1.5 - ≤ 2 x ULN	>2 x ULN	-		
Thrombosis/embolism is grade	ed in the CARDIO	VASCULAR (GENERAL) categ	gory.				
Thrombotic microangiopathy (e.g., thrombotic thrombocytopenic purpura/TTP or hemolytic uremic syndrome/HUS)	absent	-	-	laboratory findings present without clinical consequences	laboratory findings and clinical consequences, (e.g., CNS hemorrhage/ bleeding or thrombosis/ embolism or renal failure) requiring therapeutic intervention		
For BMT:	-	evidence of RBC destruction (schistocytosis) without clinical consequences	evidence of RBC destruction with elevated creatinine (≤3 x ULN)	evidence of RBC destruction with creatinine (>3 x ULN) not requiring dialysis	evidence of RBC destruction with renal failure requiring dialysis and/or encephalopathy		
Also consider Hemoglobin (H	gb), Platelets, Crea	tinine.					
Note: Must have microangiopa	thic changes on bl	ood smear (e.g., schistocytes, he	elmet cells, red cell fragment	s).			
Coagulation-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling		
		CONSTITUTION	NAL SYMPTOMS	S			
Fatigue (lethargy, malaise, asthenia)	none	increased fatigue over baseline, but not altering normal activities	moderate (e.g., decrease in performance status by 1 ECOG level or 20% Karnofsky or Lansky) or causing difficulty performing some activities	severe (e.g., decrease in performance status by ≥2 ECOG levels or 40% Karnofsky or <i>Lansky</i>) or loss of ability to perform some activities	bedridden or disabling		

	Grade								
Toxicity	0	1	2	3	4				
Fever (in the absence of neutropenia, where neutropenia is defined as AGC < 1.0 x 10 ⁹ /L)	none	38.0 - 39.0°C (100.4 - 102.2°F)	39.1 - 40.0°C (102.3 - 104.0°F)	> 40.0°C (>104.0°F) for < 24hrs	> 40.0°C (>104.0°F) for > 24hrs				
Also consider Allergic reacti	ion/hypersensitivity	•							
Note: The temperature measu	urements listed abo	ve are oral or tympanic.							
Hot flashes/flushes are grade	ed in the ENDOCRI	NE category.							
Rigors, chills	none	mild, requiring symptomatic treatment (e.g., blanket) or non- narcotic medication	severe and/or prolonged, requiring narcotic medication	not responsive to narcotic medication	-				
Sweating (diaphoresis)	normal	mild and occasional	frequent or drenching	-	-				
Weight gain	< 5%	5 - <10%	10 - <20%	≥ 20%	-				
Also consider Ascites, Edem	a, Pleural effusion.								
Weight gain - veno- occlusive disease (VOD)									
Note: The following criteria	is to be used ONLY	for weight gain associated with	Veno-Occlusive Disease.						
	<2%	≥2 - <5%	≥5 - <10%	≥10% or as ascities	≥10% or fluid retention resulting in pulmonary failure				
				>200/					
Weight loss	< 5%	5 - <10%	10 - <20%	≥20%	-				
Weight loss Also consider Vomiting, Del			10 - <20%	≥20%	-				
C			10 - <20% moderate	severe	life-threatening or disabling				
Also consider Vomiting, Del Constitutional Symptoms- Other	nydration, Diarrhea	mild							
Also consider Vomiting, Del Constitutional Symptoms- Other (Specify,)	nydration, Diarrhea	mild	moderate						
Also consider Vomiting, Del Constitutional Symptoms- Other	nydration, Diarrhea none	mild DERMATO	moderate	severe	disabling				
Also consider Vomiting, Def Constitutional Symptoms- Other (Specify,) Alopecia Bruising (in absence of grade 3 or 4 thrombocytopenia) Note: Bruising resulting from	none normal none m grade 3 or 4 thro	mild DERMATO mild hair loss localized or in	moderate PLOGY/SKIN pronounced hair loss generalized niae/purpura and Hemorrhage	severe - -	disabling				

	Grade							
Toxicity	0	1	2	3	4			
Dry skin	normal	controlled with emollients	not controlled with emollients	-	-			
Erythema multiforme (e.g., Stevens-Johnson syndrome, toxic epidermal necrolysis)	absent	-	scattered, but not generalized eruption	severe or requiring IV fluids (e.g., generalized rash or painful stomatitis)	life-threatening (e.g., exfoliative or ulcerating dermatitis or requiring enteral or parenteral nutritional support)			
Flushing	absent	present	-	-	-			
Hand-foot skin reaction	none	skin changes or dermatitis without pain (e.g., erythema, peeling)	skin changes with pain, not interfering with function	skin changes with pain, interfering with function	-			
Injection site reaction	none	pain or itching or erythema	pain or swelling, with inflammation or phlebitis	ulceration or necrosis that is severe or prolonged, or requiring surgery	-			
Nail changes	normal	discoloration or ridging (koilonychia) or pitting	partial or complete loss of nail(s) or pain in nailbeds	-	-			
Petechiae is graded in the HEI	MORRHAGE category.							
Photosensitivity	none	painless erythema	painful erythema	erythema with desquamation	-			
Pigmentation changes (e.g., vitiligo)	none	localized pigmentation changes	generalized pigmentation changes	-	-			
Pruritus	none	mild or localized, relieved spontaneously or by local measures	intense or widespread, relieved spontaneously or by systemic measures	intense or widespread and poorly controlled despite treatment	-			
Purpura is graded in the HEM	ORRHAGE category.							
Radiation dermatitis	none	faint erythema or dry desquamation	moderate to brisk erythema or a patchy moist desquamation, mostly confined to skin folds and creases; moderate edema	confluent moist desquamation, ≥1.5 cm diameter, not confined to skin folds; pitting edema	skin necrosis or ulceration of full thickness dermis; may include bleeding not induced by minor trauma or abrasion			
Note: Pain associated with rac	liation dermatitis is grad	ded separately in the PAIN c	ategory as Pain due to radiat	tion.				
Radiation recall reaction (reaction following chemotherapy in the absence of additional radiation therapy that occurs in a previous radiation port)	none	faint erythema or dry desquamation	moderate to brisk erythema or a patchy moist desquamation, mostly confined to skin folds and creases; moderate edema	confluent moist desquamation, ≥1.5 cm diameter, not confined to skin folds; pitting edema	skin necrosis or ulceration of full thickness dermis; may include bleeding not induced by minor trauma or abrasion			

		Gr	ade		
Toxicity	0	1	2	3	4
Rash/desquamation	none	macular or papular eruption or erythema without associated symptoms	macular or papular eruption or erythema with pruritus or other associated symptoms covering <50% of body surface or localized desquamation or other lesions covering <50% of body surface area	symptomatic generalized erythroderma or macular, papular or vesicular eruption or desquamation covering ≥50% of body surface area	generalized exfoliative dermatitis or ulcerative dermatitis
For BMT:	none	macular or papular eruption or erythema covering <25% of body surface area without associated symptoms	macular or papular eruption or erythema with pruritis or other associated symptoms covering ≥25 - <50% of body surface or localized desquamation or other lesions covering ≥25 - <50% of body surface area	symptomatic generalized erythroderma or symptomatic macular, papular or vesicular eruption, with bullous formation, or desquamation covering ≥50% of body surface area	generalized exfoliative dermatitis or ulcerative dermatitis or bullous formation
Also consider Allergic reaction	n/hypersensitivity.				
Note: Erythema multiforme (S	tevens-Johnson syndro	me) is graded separately as l	Erythema multiforme.		
Urticaria (hives, welts, wheals)	none	requiring no medication	requiring PO or topical treatment or IV medication or steroids for <24 hours	requiring IV medication or steroids for ≥24 hours	-
Wound- infectious	none	cellulitis	superficial infection	infection requiring IV antibiotics	necrotizing fascitis
Wound- non-infectious	none	incisional separation	incisional hernia	fascial disruption without evisceration	fascial disruption with evisceration
Dermatology/Skin-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		ENDO	CRINE		
Cushingoid appearance (e.g., moon face with or without buffalo hump, centripetal obesity, cutaneous striae)	absent	-	present	-	-
Also consider Hyperglycemia,	••				
Feminization of male	absent	-	-	present	-
Gynecomastia	none	mild	pronounced or painful	pronounced or painful and requiring surgery	-
Hot flashes/flushes	none	mild or no more than 1 per day	moderate and greater than 1 per day	-	-
Hypothyroidism	absent	asymptomatic,TSH elevated, no therapy given	symptomatic or thyroid replacement treatment given	patient hospitalized for manifestations of hypothyroidism	myxedema coma
Masculinization of female	absent	-	-	present	-

		Gr	ade		
Toxicity	0	1	2	3	4
SIADH (syndrome of inappropriate antidiuretic hormone)	absent	-	-	present	-
Endocrine-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		GASTROIN	NTESTINAL		
Amylase is graded in the MET	TABOLIC/LABORA	ΓORY category.			
Anorexia	none	loss of appetite	oral intake significantly decreased	requiring IV fluids	requiring feeding tube or parenteral nutrition
Ascites (non-malignant)	none	asymptomatic	symptomatic, requiring diuretics	symptomatic, requiring therapeutic paracentesis	life-threatening physiologic consequences
Colitis	none	-	abdominal pain with mucus and/or blood in stool	abdominal pain, fever, change in bowel habits with ileus or peritoneal signs, and radiographic or biopsy documentation	perforation or requiring surgery or toxic megacolon
Also consider Hemorrhage/ble Rectal bleeding/hematochezia Constipation		requiring stool softener	hage/bleeding without grade requiring laxatives	obstipation requiring	obstruction or toxic
		or dietary modification		manual evacuation or enema	megacolon
Dehydration	none	dry mucous membranes and/or diminished skin turgor	requiring IV fluid replacement (brief)	requiring IV fluid replacement (sustained)	physiologic consequences requiring intensive care; hemodynamic collapse
Also consider Hypotension, D	iarrhea, Vomiting, St	omatitis/pharyngitis (oral/phar	ryngeal mucositis).		
Diarrhea Patients without colostomy:	none	increase of < 4 stools/day over pre- treatment	increase of 4-6 stools/day, or nocturnal stools	increase of ≥7 stools/day or incontinence; or need for parenteral support for dehydration	physiologic consequences requiring intensive care; or hemodynamic collapse
Patients with a colostomy:	none	mild increase in loose, watery colostomy output compared with pretreatment	moderate increase in loose, watery colostomy output compared with pretreatment, but not interfering with normal activity	severe increase in loose, watery colostomy output compared with pretreatment, interfering with normal activity	physiologic consequences, requiring intensive care; or hemodynamic collapse
For BMT	none	>500 - ≤1000ml of diarrhea/day	>1000 - ≤1500ml of diarrhea/day	>1500ml of diarrhea/day	severe abdominal pain with or without ileus
For Pediatric BMT:		>5 - £10 ml/kg of diarrhea/day	>10 - £15 ml/kg of diarrhea/day	>15 ml/kg of diarrhea/day	-
Also consider Hemorrhage/ble Hypotension.	eeding with grade 3 or	r 4 thrombocytopenia, Hemorr	hage/bleeding without grade	e 3 or 4 thrombocytopenia, F	Pain, Dehydration,

Grade								
Toxicity	0	1	2	3	4			
Duodenal ulcer (requires radiographic or endoscopic documentation)	none	-	requiring medical management or non- surgical treatment	uncontrolled by outpatient medical management; requiring hospitalization	perforation or bleeding, requiring emergency surgery			
Dyspepsia/heartburn	none	mild	moderate	severe	-			
Dysphagia, esophagitis, odynophagia (painful swallowing)	none	mild dysphagia, but can eat regular diet	dysphagia, requiring predominantly pureed, soft, or liquid diet	dysphagia, requiring IV hydration	complete obstruction (cannot swallow saliva) requiring enteral or parenteral nutritional support, or perforation			
Note: If toxicity is radiation-r	elated, grade <u>either</u> und	er Dysphagia- esophageal re	lated to radiation or Dyspha	gia- pharyngeal related to ra	diation.			
Dysphagia- <u>esophageal</u> related to radiation	none	mild dysphagia, but can eat regular diet	dysphagia, requiring predominantly liquid, pureed or soft diet	dysphagia requiring feeding tube, IV hydration or hyperalimentation	complete obstruction (cannot swallow saliva); ulceration with bleeding not induced by minor trauma or abrasion or perforation			
Also consider Pain due to rad	iation, Mucositis due to	radiation.						
Note: Fistula is graded separa	tely as Fistula- esophag	eal.						
Dysphagia - <u>pharyngeal</u> related to radiation	none	mild dysphagia, but can eat regular diet	dysphagia, requiring predominantly pureed, soft, or liquid diet	dysphagia, requiring feeding tube, IV hydration or hyperalimentation	complete obstruction (cannot swallow saliva); ulceration with bleeding not induced by minor trauma or abrasion or perforation			
Also consider Pain due to rad								
Note: Fistula is graded separa					ma avidina avimaami			
Fistula- esophageal	none	-	-	present	requiring surgery			
Fistula- intestinal	none	<u>-</u>	-	present	requiring surgery			
Fistula- pharyngeal	none	-	-	present	requiring surgery			
Fistula- rectal/anal	none	-	-	present	requiring surgery			
Flatulence	none	mild	moderate	-	-			
Gastric ulcer (requires radiographic or endoscopic documentation)	none	-	requiring medical management or non- surgical treatment	bleeding without perforation, uncontrolled by outpatient medical management; requiring hospitalization or surgery	perforation or bleeding, requiring emergency surgery			
(requires radiographic or			management or non- surgical treatment	perforation, uncontrolled by outpatient medical management; requiring hospitalization or surgery	requiring emergency			
(requires radiographic or endoscopic documentation)			management or non- surgical treatment	perforation, uncontrolled by outpatient medical management; requiring hospitalization or surgery	requiring emergency			

	Grade								
Toxicity	0	1	2	3	4				
Hematemesis is graded in	the HEMORRHAGE	E category.							
Hematochezia is graded in	the HEMORRHAG	E category as Rectal bleeding/hem	natochezia.						
Ileus (or neuroconstipation	n) none	-	intermittent, not requiring intervention	requiring non-surgical intervention	requiring surgery				
Mouth dryness	normal	mild	moderate	-	-				
Mucositis									
		in the GASTROINTESTINAL cattis; or the RENAL/GENITOURIN			tomatitis/pharyngitis				
Radiation-related n	nucositis is graded as	Mucositis due to radiation.							
Mucositis due to radiation	none	erythema of the mucosa	patchy pseudomembranous reaction (patches generally ≤ 1.5 cm in diameter and non- contiguous)	confluent pseudomembranous reaction (contiguous patches generally > 1.5 cm in diameter)	necrosis or deep ulceration; may include bleeding not induced by minor trauma or abrasion				
Also consider Pain due to	radiation.								
Note: Grade radiation mu	acositis of the larynx	here.							
Dysphagia related on the site of treatm		aded as <u>either</u> Dysphagia- esophag	real related to radiation or D	ysphagia- pharyngeal related	to radiation, depending				
Nausea	none	able to eat	oral intake significantly decreased	no significant intake, requiring IV fluids	-				
Pancreatitis	none	-	-	abdominal pain with pancreatic enzyme elevation	complicated by shock (acute circulatory failure)				
Also consider Hypotensio	n.								
Note: Asymptomatic amy	lase and Amylase are	graded in the METABOLIC/LAB	ORATORY category.						
Pharyngitis is graded in th	ne GASTROINTEST	INAL category as Stomatitis/phary	ngitis (oral/pharyngeal muc	ositis).					
Proctitis	none	increased stool frequency, occasional blood-streaked stools, or rectal discomfort (including hemorrhoids), not requiring medication	increased stool frequency, bleeding, mucus discharge, or rectal discomfort requiring medication; anal fissure	increased stool frequency/diarrhea, requiring parenteral support; rectal bleeding, requiring transfusion; or persistent mucus discharge, necessitating pads	perforation, bleeding or necrosis or other life- threatening complication requiring surgical intervention (e.g., colostomy)				
Also consider Hemorrhage	e/bleeding with grade	e 3 or 4 thrombocytopenia, Hemori	rhage/bleeding without grad	e 3 or 4 thrombocytopenia, a	nd Pain due to radiation.				
Note: Fistula is graded se	eparately as Fistula- re	ectal/anal.							
Proctitis occurring Appendix IV)	more than 90 days af	ter the start of radiation therapy is	graded in the RTOG/EORT	C Late Radiation Morbidity	Scoring Scheme. (See				
Salivary gland changes	none	slightly thickened saliva/may have slightly altered taste (e.g., metallic); additional fluids may be required	thick, ropy, sticky saliva; markedly altered taste; alteration in diet required	-	acute salivary gland necrosis				

		Gr	rade		
Toxicity	0	1	2	3	4
Sense of smell	normal	slightly altered	markedly altered	-	-
Stomatitis/pharyngitis (oral/pharyngeal mucositis)	none	painless ulcers, erythema, or mild soreness in the absence of lesions	painful erythema, edema, or ulcers, but can eat or swallow	painful erythema, edema, or ulcers requiring IV hydration	severe ulceration or requires parenteral or enteral nutritional support or prophylatic intubation
For BMT:	none	painless ulcers, erythema, or mild soreness in the absence of lesions	painful erythema, edema or ulcers but can swallow	painful erythema, edema, or ulcers preventing swallowing or requiring hydration or parenteral (or enteral) nutritional support	severe ulceration requiring prophylactic intubation or resulting in documented aspiration pneumonia
Note: Radiation-related mucos	sitis is graded as Mucos	itis due to radiation.			
Taste disturbance (dysgeusia)	normal	slightly altered	markedly altered	-	-
Typhlitis (inflammation of the cecum)	none	-	-	abdominal pain, diarrhea, fever, or radiographic documentation	perforation, bleeding or necrosis or other life- threatening complication requiring surgical intervention (e.g., colostomy)
Also consider Hemorrhage/ble Febrile/neutropenia.	eeding with grade 3 or 4	thrombocytopenia, Hemorr	hage/bleeding without grade	e 3 or 4 thrombocytopenia, I	Hypotension,
Vomiting	none	1 episode in 24 hours over pretreatment	2-5 episodes in 24 hours over pretreatment	≥6 episodes in 24 hours over pretreatment; or need for IV fluids	Requiring parenteral nutrition; or physiologic consequences requiring intensive care; hemodynamic collapse
Also consider Dehydration.					
Weight gain is graded in the C	CONSTITUTIONAL SY	MPTOMS category.			
Weight loss is graded in the C	ONSTITUTIONAL SY	MPTOMS category.			
Gastrointestinal-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		HEMOD	DHACE		

HEMORRHAGE

Note: Transfusion in this section refers to pRBC infusion.

For <u>any</u> bleeding with grade 3 or 4 platelets (< 50,000), <u>always</u> grade Hemorrhage/bleeding with grade 3 or 4 thrombocytopenia. Also consider platelets, transfusion-pRBCS, and transfusion-platelets in addition to the grade that incorporates the site or type of bleeding.

If the site or type of hemorrhage/bleeding is listed, also use the grading that incorporates the site of bleeding: CNS hemorrhage/bleeding, Hematuria, Hematemesis, Hemoptysis, Hemorrhage/bleeding with surgery, Melena/lower GI bleeding, Petechiae/purpura (Hemorrhage/bleeding into skin), Rectal bleeding/hematochezia, Vaginal bleeding.

If the platelet count is \geq 50,000 and the site or type of bleeding is listed, grade the specific site. If the site or type is <u>not</u> listed and the platelet count is \geq 50,000, grade Hemorrhage/bleeding without grade 3 or 4 thrombocytopenia and specify the site or type in the OTHER category.

		G	rade		
Toxicity	0	1	2	3	4
Hemorrhage/bleeding with grade 3 or 4 thrombocytopenia	none	mild without transfusion		requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Also consider Platelets, Hemo	oglobin, Transfusion-	platelet, Transfusion-pRBCs.			
Note: This toxicity must be g listed, grade as Other i			ytopenia. Also grade the site	or type of hemorrhage/bleed	ling. If the site is not
Hemorrhage/bleeding without grade 3 or 4 thrombocytopenia	none	mild without transfusion		requiring transfusion	catastrophic bleeding requiring major non- elective intervention
Also consider Platelets, Hemo	oglobin, Transfusion-	platelet, Transfusion-pRBCs.			
Note: Bleeding in the absence HEMORRHAGE cates		ombocytopenia is graded here Other in the HEMORRHAGE		be of bleeding is not listed els	sewhere in the
CNS hemorrhage/bleeding	none	-	-	bleeding noted on CT or other scan with no clinical consequences	hemorrhagic stroke or hemorrhagic vascular event (CVA) with neurologic signs and symptoms
Epistaxis	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Hematemesis	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Hematuria (in the absence of vaginal bleeding)	none	microscopic only	intermittent gross bleeding, no clots	persistent gross bleeding or clots; may require catheterization or instrumentation, or transfusion	open surgery or necrosis or deep bladder ulceration
Hemoptysis	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Hemorrhage/bleeding associated with surgery	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Note: Expected blood loss at t		•			
Melena/GI bleeding	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention
Petechiae/purpura (hemorrhage/bleeding into skin or mucosa)	none	rare petechiae of skin	petechiae or purpura in dependent areas of skin	generalized petechiae or purpura of skin or petechiae of any mucosal site	-

Grade									
Toxicity	0	1	2	3	4				
Rectal bleeding/ hematochezia	none	mild without transfusion or medication	persistent, requiring medication (e.g., steroid suppositories) and/or break from radiation treatment	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention				
Vaginal bleeding	none	spotting, requiring < 2 pads per day	requiring ≥ 2 pads per day, but not requiring transfusion	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention				
Hemorrhage-Other (Specify site,)	none	mild without transfusion	-	requiring transfusion	catastrophic bleeding, requiring major non- elective intervention				
		HEF	PATIC						
Alkaline phosphatase	WNL	> ULN - 2.5 x ULN	> 2.5 - 5.0 x ULN	> 5.0 - 20.0 x ULN	> 20.0 x ULN				
Bilirubin	WNL	> ULN - 1.5 x ULN	> 1.5 - 3.0 x ULN	> 3.0 - 10.0 x ULN	> 10.0 x ULN				
Bilirubin- graft versus host dis	sease (GVHD)								
Note: The following criteria ar	re used only for bi	lirubin associated with graft vers	sus host disease.						
	normal	≥2 - <3 mg/100 ml	≥3 - <6 mg/100 ml	≥6 - <15 mg/100 ml	≥15 mg/100 ml				
GGT (γ - Glutamyl transpeptidase)	WNL	> ULN - 2.5 x ULN	> 2.5 - 5.0 x ULN	> 5.0 - 20.0 x ULN	> 20.0 x ULN				
Hepatic enlargement	absent	-	-	present	-				
Note: Grade Hepatic enlargem	nent only for chang	ges related to VOD or other treat	ment related toxicity.						
Hypoalbuminemia	WNL	<lln -="" 3="" dl<="" g="" td=""><td>≥2 - <3 g/dl</td><td><2 g/dl</td><td>-</td></lln>	≥2 - <3 g/dl	<2 g/dl	-				
Liver dysfunction/failure (clinical)	normal	-	-	asterixis	encephalopathy or coma				
Note: Documented viral hepat	itis is graded in the	e INFECTION category.							
Portal vein flow	normal	-	decreased portal vein flow	reversal/retrograde portal vein flow	-				
SGOT (AST) (serum glutamic oxaloacetic transaminase)	WNL	> ULN - 2.5 x ULN	> 2.5 - 5.0 x ULN	> 5.0 - 20.0 x ULN	> 20.0 x ULN				
SGPT (ALT) (serum glutamic pyruvic transaminase)	WNL	> ULN - 2.5 x ULN	> 2.5 - 5.0 x ULN	> 5.0 - 20.0 x ULN	> 20.0 x ULN				
Hepatic-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling				
		INFECTION/FEBR	ILE NEUTROPE	NIA					
Catheter-related infection	none	mild, no active treatment	moderate, localized infection, requiring local or oral treatment	severe, systemic infection, requiring IV antibiotic or antifungal treatment or hospitalization	life-threatening sepsis (e.g., septic shock)				

	Grade								
Toxicity	0	1	2	3	4				
Febrile neutropenia (fever of unknown origin without clinically or microbiologically documented infection)	none	-	-	Present	Life-threatening sepsis (e.g., septic shock)				
(ANC < 1.0×10^9 /L, fever ≥ 38.5 °C)									
Note: Hypothermia instead o	f fever may be ass	ociated with neutropenia and is g	raded here.						
Infection (documented clinically or microbiologically) with grade 3 or 4 neutropenia	none	-	-	present	life-threatening sepsis (e.g., septic shock)				
$(ANC < 1.0 \times 10^9/L)$									
Note: Hypothermia instead of grade as Febrile neutro		sociated with neutropenia and is a	graded here. In the absence o	f documented infection with	grade 3 or 4 neutropenia,				
Infection with unknown ANC	none	-	-	present	life-threatening sepsis (e.g., septic shock)				
Note: This toxicity criterion i	s used in the rare	case when ANC is unknown.							
Infection without neutropenia	none	mild, no active treatment	moderate, localized infection, requiring local or oral treatment	severe, systemic infection, requiring IV antibiotic or antifungal treatment, or hospitalization	life-threatening sepsis (e.g., septic shock)				
Infection/Febrile Neutropenia-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling				
Wound-infectious is graded i	n the DERMATO	LOGY/SKIN category.							
		LYMP	HATICS						
Lymphatics	normal	mild lymphedema	moderate lymphedema requiring compression; lymphocyst	severe lymphedema limiting function; lymphocyst requiring surgery	severe lymphedema limiting function with ulceration				
Lymphatics-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling				
		METABOLIC	LABORATORY						
Acidosis (metabolic or respiratory)	normal	pH < normal, but ≥7.3	-	pH < 7.3	pH < 7.3 with life- threatening physiologic consequences				
Alkalosis (metabolic or respiratory)	normal	pH > normal, but ≤7.5	-	pH > 7.5	pH > 7.5 with life- threatening physiologic consequences				
Amylase	WNL	> ULN - 1.5 x ULN	> 1.5 - 2.0 x ULN	> 2.0 - 5.0 x ULN	>5.0 x ULN				
•									

		Gr	rade		
Toxicity	0	1	2	3	4
CPK (creatine phosphokinase)	WNL	> ULN - 2.5 x ULN	> 2.5 - 5 x ULN	> 5 - 10 x ULN	> 10 x ULN
Hypercalcemia	WNL	> ULN - 11.5 mg/dl > ULN - 2.9 mmol/L	>11.5 - 12.5 mg/dl > 2.9 - 3.1 mmol/L	>12.5 - 13.5 mg/dl > 3.1 - 3.4 mmol/L	> 13.5 mg/dl > 3.4 mmol/L
Hypercholesterolemia	WNL	> ULN - 300 mg/dl > ULN - 7.75 mmol/L	> 300 - 400 mg/dl > 7.75 - 10.34 mmol/L	> 400 - 500 mg/dl >10.34 - 12.92 mmol/L	> 500 mg/dl > 12.92 mmol/L
Hyperglycemia	WNL	> ULN - 160 mg/dl > ULN - 8.9 mmol/L	> 160 - 250 mg/dl > 8.9 - 13.9 mmol/L	> 250 - 500 mg/dl > 13.9 - 27.8 mmol/L	> 500 mg/dl > 27.8 mmol/L or ketoacidosis
Hyperkalemia	WNL	> ULN - 5.5 mmol/L	> 5.5 - 6.0 mmol/L	> 6.0 - 7.0 mmol/L	> 7.0 mmol/L
Hypermagnesemia	WNL	> ULN - 3.0 mg/dl > ULN - 1.23 mmol/L	-	> 3.0 - 8.0 mg/dl > 1.23 - 3.30 mmol/L	> 8.0 mg/dl > 3.30 mmol/L
Hypernatremia	WNL	> ULN - 150 mmol/L	>150 - 155 mmol/L	>155 - 160 mmol/L	>160 mmol/L
Hypertriglyceridemia	WNL	> ULN - 2.5 x ULN	> 2.5 - 5.0 x ULN	> 5.0 - 10 x ULN	> 10 x ULN
Hyperuricemia	WNL	> ULN - ≤ 10 mg/dl ≤ 0.59 mmol/L without physiologic consequences	-	> ULN - ≤ 10 mg/dl ≤ 0.59 mmol/L with physiologic consequences	> 10 mg/dl > 0.59 mmol/L
Also consider Tumor lysis syr	drome, Renal failure,	Creatinine, Potassium.			
Hypocalcemia	WNL	<lln -="" 8.0="" dl<br="" mg=""><lln -="" 2.0="" l<="" mmol="" td=""><td>7.0 - < 8.0 mg/dl 1.75 - < 2.0 mmol/L</td><td>6.0 - < 7.0 mg/dl 1.5 - < 1.75 mmol/L</td><td><6.0 mg/dl < 1.5 mmol/L</td></lln></lln>	7.0 - < 8.0 mg/dl 1.75 - < 2.0 mmol/L	6.0 - < 7.0 mg/dl 1.5 - < 1.75 mmol/L	<6.0 mg/dl < 1.5 mmol/L
Hypoglycemia	WNL	<lln -="" 55="" dl<br="" mg=""><lln -="" 3.0="" l<="" mmol="" td=""><td>40 - < 55 mg/dl 2.2 - < 3.0 mmol/L</td><td>30 - < 40 mg/dl 1.7 - < 2.2 mmol/L</td><td>< 30 mg/dl < 1.7 mmol/L</td></lln></lln>	40 - < 55 mg/dl 2.2 - < 3.0 mmol/L	30 - < 40 mg/dl 1.7 - < 2.2 mmol/L	< 30 mg/dl < 1.7 mmol/L
Hypokalemia	WNL	<lln -="" 3.0="" l<="" mmol="" td=""><td>-</td><td>2.5 - <3.0 mmol/L</td><td><2.5 mmol/L</td></lln>	-	2.5 - <3.0 mmol/L	<2.5 mmol/L
Hypomagnesemia	WNL	<lln -="" 1.2="" dl<br="" mg=""><lln -="" 0.5="" l<="" mmol="" td=""><td>0.9 - <1.2 mg/dl 0.4 - < 0.5 mmol/L</td><td>0.7 - < 0.9 mg/dl 0.3 - < 0.4 mmol/L</td><td>< 0.7 mg/dl < 0.3 mmol/L</td></lln></lln>	0.9 - <1.2 mg/dl 0.4 - < 0.5 mmol/L	0.7 - < 0.9 mg/dl 0.3 - < 0.4 mmol/L	< 0.7 mg/dl < 0.3 mmol/L
Hyponatremia	WNL	<lln -="" 130="" l<="" mmol="" td=""><td>-</td><td>120 - <130 mmol/L</td><td><120 mmol/L</td></lln>	-	120 - <130 mmol/L	<120 mmol/L
Hypophosphatemia	WNL	<lln -2.5="" dl<br="" mg=""><lln -="" 0.8="" l<="" mmol="" td=""><td>≥2.0 - <2.5 mg/dl ≥0.6 - <0.8 mmol/L</td><td>≥1.0 - <2.0 mg/dl ≥0.3 - <0.6 mmol/L</td><td>< 1.0 mg/dl <0.3 mmol/L</td></lln></lln>	≥2.0 - <2.5 mg/dl ≥0.6 - <0.8 mmol/L	≥1.0 - <2.0 mg/dl ≥0.3 - <0.6 mmol/L	< 1.0 mg/dl <0.3 mmol/L
Hypothyroidism is graded in t	he ENDOCRINE cate	gory.			
Lipase	WNL	> ULN - 1.5 x ULN	> 1.5 - 2.0 x ULN	> 2.0 - 5.0 x ULN	> 5.0 x ULN
Metabolic/Laboratory-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		MUSCULO	SKELETAL		
Arthralgia is graded in the PA	IN category.				
Arthritis	none	mild pain with inflammation, erythema or joint swelling but not interfering with function	moderate pain with inflammation, erythema, or joint swelling interfering with function, but not interfering with activities of daily living	severe pain with inflammation, erythema, or joint swelling and interfering with activities of daily living	disabling

		Gr	ade		
Toxicity	0	1	2	3	4
Muscle weakness (not due to neuropathy)	normal	asymptomatic with weakness on physical exam	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	bedridden or disabling
Myalgia is graded in the PAIN	I category.				
Myositis (inflammation/damage of muscle)	none	mild pain, not interfering with function	pain interfering with function, but not interfering with activities of daily living	pain interfering with function and interfering with activities of daily living	bedridden or disabling
Also consider CPK.					
Note: Myositis implies muscle	e damage (i.e., elev	rated CPK).			
Osteonecrosis (avascular necrosis)	none	asymptomatic and detected by imaging only	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	symptomatic; or disabling
Musculoskeletal-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		NEUR (OLOGY		
Aphasia, receptive and/or exp	ressive, is graded u	ander Speech impairment in the N	EUROLOGY category.		
Arachnoiditis/meningismus/radiculitis	absent	mild pain not interfering with function	moderate pain interfering with function, but not interfering with activities of daily living	severe pain interfering with activities of daily living	unable to function or perform activities of daily living; bedridden paraplegia
Also consider Headache, Von	niting, Fever.				
Ataxia (incoordination)	normal	asymptomatic but abnormal on physical exam, and not interfering with function	mild symptoms interfering with function, but not interfering with activities of daily living	moderate symptoms interfering with activities of daily living	bedridden or disabling
CNS cerebrovascular ischemia	none	-	-	transient ischemic event or attack (TIA)	permanent event (e.g., cerebral vascular accident)
CNS hemorrhage/bleeding is §	graded in the HEM	ORRHAGE category.			
Cognitive disturbance/ learning problems	none	cognitive disability; not interfering with work/school performance; preservation of intelligence	cognitive disability; interfering with work/school performance; decline of 1 SD (Standard Deviation) or loss of developmental milestones	cognitive disability; resulting in significant impairment of work/school performance; cognitive decline > 2 SD	inability to work/frank mental retardation

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Grade									
Toxicity	0	1	2	3	4				
Confusion	normal	confusion or disorientation or attention deficit of brief duration; resolves spontaneously with no sequelae	confusion or disorientation or attention deficit interfering with function, but not interfering with activities of daily living	confusion or delirium interfering with activities of daily living	harmful to others or self; requiring hospitalization				
Cranial neuropathy is graded	in the NEUROLO	GY category as Neuropathy-crani	al.						
Delusions	normal	-	-	present	toxic psychosis				
Depressed level of consciousness	normal	somnolence or sedation not interfering with function	somnolence or sedation interfering with function, but not interfering with activities of daily living	obtundation or stupor; difficult to arouse; interfering with activities of daily living	coma				
Note: Syncope (fainting) is g	raded in the NEUF	ROLOGY category.							
Dizziness/lightheadedness	none	not interfering with function	interfering with function, but not interfering with activities of daily living	interfering with activities of daily living	bedridden or disabling				
Dysphasia, receptive and/or e	expressive, is grade	ed under Speech impairment in the	e NEUROLOGY category.						
Extrapyramidal/ involuntary movement/ restlessness	none	mild involuntary movements not interfering with function	moderate involuntary movements interfering with function, but not interfering with activities of daily living	severe involuntary movements or torticollis interfering with activities of daily living	bedridden or disabling				
Hallucinations	normal	-	-	present	toxic psychosis				
Headache is graded in the PA	AIN category.								
Insomnia	normal	occasional difficulty sleeping not interfering with function	difficulty sleeping interfering with function, but not interfering with activities of daily living	frequent difficulty sleeping, interfering with activities of daily living	-				
Note: This toxicity is graded	when insomnia is	related to treatment. If pain or other	er symptoms interfere with s	leep do NOT grade as inson	nnia.				
Irritability (children <3 years of age)	normal	mild; easily consolable	moderate; requiring increased attention	severe; inconsolable	-				
Leukoencephalopathy associated radiological findings	none	mild increase in SAS (subarachnoid space) and/or mild ventriculomegaly; and/or small (+/- multiple) focal T2 hyperintensities, involving periventricular white matter or < 1/3 of susceptible areas of cerebrum	moderate increase in SAS; and/or moderate ventriculomegaly; and/or focal T2 hyperintensities extending into centrum ovale; or involving 1/3 to 2/3 of susceptible areas of cerebrum	severe increase in SAS; severe ventriculomegaly; near total white matter T2 hyperintensities or diffuse low attenuation (CT); focal white matter necrosis (cystic)	severe increase in SAS; severe ventriculomegaly; diffuse low attenuation with calcification (CT); diffuse white matter necrosis (MRI)				
Memory loss	normal	memory loss not interfering with function	memory loss interfering with function, but not interfering with activities of daily living	memory loss interfering with activities of daily living	amnesia				

		Gr	ade		
Toxicity	0	1	2	3	4
Mood alteration- anxiety agitation	normal	mild mood alteration not interfering with function	moderate mood alteration interfering with function, but not interfering with activities of daily living	severe mood alteration interfering with activities of daily living	suicidal ideation or danger to self
Mood alteration- depression	normal	mild mood alteration not interfering with function	moderate mood alteration interfering with function, but not interfering with activities of daily living	severe mood alteration interfering with activities of daily living	suicidal ideation or danger to self
Mood alteration- euphoria	normal	mild mood alteration not interfering with function	moderate mood alteration interfering with function, but not interfering with activities of daily living	severe mood alteration interfering with activities of daily living	danger to self
Neuropathic pain is graded in	the PAIN category.				
Neuropathy- cranial	absent	-	present, not interfering with activities of daily living	present, interfering with activities of daily living	life-threatening, disabling
Neuropathy- motor	normal	subjective weakness but no objective findings	mild objective weakness interfering with function, but not interfering with activities of daily living	objective weakness interfering with activities of daily living	paralysis
Neuropathy-sensory	normal	loss of deep tendon reflexes or paresthesia (including tingling) but not interfering with function	objective sensory loss or paresthesia (including tingling), interfering with function, but not interfering with activities of daily living	sensory loss or paresthesia interfering with activities of daily living	permanent sensory loss that interferes with function
Nystagmus	absent	present	-	-	-
Also consider Vision-double v	vision.				
Personality/behavioral	normal	change, but not disruptive to patient or family	disruptive to patient or family	disruptive to patient and family; requiring mental health intervention	harmful to others or self; requiring hospitalization
Pyramidal tract dysfunction (e.g., ↑ tone, hyperreflexia, positive Babinski, ↓ fine motor coordination)	normal	asymptomatic with abnormality on physical examination	symptomatic or interfering with function but not interfering with activities of daily living	interfering with activities of daily living	bedridden or disabling; paralysis
Seizure(s)	none	-	seizure(s) self-limited and consciousness is preserved	seizure(s) in which consciousness is altered	seizures of any type which are prolonged, repetitive, or difficult to control (e.g., status epilepticus, intractable epilepsy)
Speech impairment (e.g., dysphasia or aphasia)	normal	-	awareness of receptive or expressive dysphasia, not impairing ability to communicate	receptive or expressive dysphasia, impairing ability to communicate	inability to communicate
	absent			present	<u> </u>

		Gr	ade		
Toxicity	0	1	2	3	4
Tremor	none	mild and brief or intermittent but not interfering with function	moderate tremor interfering with function, but not interfering with activities of daily living	severe tremor interfering with activities of daily living	-
Vertigo	none	not interfering with function	interfering with function, but not interfering with activities of daily living	interfering with activities of daily living	bedridden or disabling
Neurology-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		OCULAR	R/VISUAL		
Cataract	none	asymptomatic	symptomatic, partial visual loss	symptomatic, visual loss requiring treatment or interfering with function	-
Conjunctivitis	none	abnormal ophthalmologic changes, but asymptomatic or symptomatic without visual impairment (i.e., pain and irritation)	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-
Dry eye	normal	mild, not requiring treatment	moderate or requiring artificial tears	-	-
Glaucoma	none	increase in intraocular pressure but no visual loss	increase in intraocular pressure with retinal changes	visual impairment	unilateral or bilateral loss of vision (blindness)
Keratitis (corneal inflammation/ corneal ulceration)	none	abnormal ophthalmologic changes but asymptomatic or symptomatic without visual impairment (i.e., pain and irritation)	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	unilateral or bilateral loss of vision (blindness)
Tearing (watery eyes)	none	mild: not interfering with function	moderate: interfering with function, but not interfering with activities of daily living	interfering with activities of daily living	-
Vision- blurred vision	normal	-	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-
Vision- double vision (diplopia)	normal	-	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-
Vision- flashing lights/floaters	normal	mild, not interfering with function	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-

		Gr	ade		
Toxicity	0	1	2	3	4
Vision- night blindness (nyctalopia)	normal	abnormal electro- retinography but asymptomatic	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-
Vision- photophobia	normal	-	symptomatic and interfering with function, but not interfering with activities of daily living	symptomatic and interfering with activities of daily living	-
Ocular/Visual-Other (Specify,)	normal	mild	moderate	severe	unilateral or bilateral loss of vision (blindness)
		PA	IN		
Abdominal pain or cramping	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Arthralgia (joint pain)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Arthritis (joint pain with clinic	al signs of inflammatio	n) is graded in the MUSCUI	LOSKELETAL category.		
Bone pain	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Chest pain (non-cardiac and non- pleuritic)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Dysmenorrhea	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Dyspareunia	none	mild pain not interfering with function	moderate pain interfering with sexual activity	severe pain preventing sexual activity	-
Dysuria is graded in the RENA	AL/GENITOURINARY	category.			
Earache (otalgia)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling
Headache	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling

Grade								
Toxicity	0	1	2	3	4			
Hepatic pain	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Myalgia (muscle pain)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Neuropathic pain (e.g., jaw pain, neurologic pain, phantom limb pain, post-infectious neuralgia, or painful neuropathies)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Pain due to radiation	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Pelvic pain	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Pleuritic pain	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Rectal or perirectal pain (proctalgia)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Tumor pain (onset or exacerbation of tumor pain due to treatment)	none	mild pain not interfering with function	moderate pain: pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain: pain or analgesics severely interfering with activities of daily living	disabling			
Tumor flair is graded in the SY	NDROME category.							
Pain-Other (Specify,)	none	mild	moderate	severe	disabling			
		PULMO	ONARY					
Adult Respiratory Distress Syndrome (ARDS)	absent		-	-	present			
Apnea	none	-	-	present	requiring intubation			

		Gr	ade		
Toxicity	0	1	2	3	4
Carbon monoxide diffusion capacity (DL_{CO})	≥ 90% of pretreatment or normal value	≥75 - <90% of pretreatment or normal value	≥50 - <75% of pretreatment or normal value	≥25 - <50% of pretreatment or normal value	< 25% of pretreatment or normal value
Cough	absent	mild, relieved by non- prescription medication	requiring narcotic antitussive	severe cough or coughing spasms, poorly controlled or unresponsive to treatment	-
Dyspnea (shortness of breath)	normal	-	dyspnea on exertion	dyspnea at normal level of activity	dyspnea at rest or requiring ventilator support
FEV ₁	≥ 90% of pretreatment or normal value	≥75 - <90% of pretreatment or normal value	≥50 - <75% of pretreatment or normal value	≥25 - <50% of pretreatment or normal value	< 25% of pretreatment or normal value
Hiccoughs (hiccups, singultus)	none	mild, not requiring treatment	moderate, requiring treatment	severe, prolonged, and refractory to treatment	-
Нурохіа	normal	-	decreased O ₂ saturation with exercise	decreased O ₂ saturation at rest, requiring supplemental oxygen	decreased O ₂ saturation, requiring pressure support (CPAP) or assisted ventilation
Pleural effusion (non-malignant)	none	asymptomatic and not requiring treatment	symptomatic, requiring diuretics	symptomatic, requiring O ₂ or therapeutic thoracentesis	life-threatening (e.g., requiring intubation)
Pleuritic pain is graded in the F	PAIN category.				
Pneumonitis/pulmonary infiltrates	none	radiographic changes but asymptomatic or symptoms not requiring steroids	radiographic changes and requiring steroids or diuretics	radiographic changes and requiring oxygen	radiographic changes and requiring assisted ventilation
Pneumothorax	none	no intervention required	chest tube required	sclerosis or surgery required	life-threatening
Pulmonary embolism is graded	l as Thrombosis/embol	ism in the CARDIOVASCU	LAR (GENERAL) category		
Pulmonary fibrosis	none	radiographic changes, but asymptomatic or symptoms not requiring steroids	requiring steroids or diuretics	requiring oxygen	requiring assisted ventilation
Note: Radiation-related pulmo	nary fibrosis is graded	in the RTOG/EORTC Late I	Radiation Morbidity Scoring	Scheme- Lung. (See Apper	ndix IV)
Voice changes/stridor/larynx (e.g., hoarseness, loss of voice, laryngitis)	normal	mild or intermittent hoarseness	persistent hoarseness, but able to vocalize; may have mild to moderate edema	whispered speech, not able to vocalize; may have marked edema	marked dyspnea/stridor requiring tracheostomy or intubation
Note: Cough from radiation is	graded as cough in the	e PULMONARY category.			
			ucositis due to radiation in the HEMORRHAGE of		category. Radiation-
Pulmonary-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling

		Gr	rade		
Toxicity	0	1	2	3	4
		RENAL/GEN	ITOURINARY		
Bladder spasms	absent	mild symptoms, not requiring intervention	symptoms requiring antispasmotic	severe symptoms requiring narcotic	-
Creatinine	WNL	> ULN - 1.5 x ULN	> 1.5 - 3.0 x ULN	> 3.0 - 6.0 x ULN	> 6.0 x ULN
Note: Adjust to age-appropria	ate levels for pediatric p	patients.			
Dysuria (painful urination)	none	mild symptoms requiring no intervention	symptoms relieved with therapy	symptoms not relieved despite therapy	Ŧ.
Fistula or GU fistula (e.g., vaginal, vesicovaginal)	none	-	-	requiring intervention	requiring surgery
Hemoglobinuria	-	present	-	-	-
Hematuria (in the absence of	vaginal bleeding) is gra	aded in the HEMORRHAGE	category.		
Incontinence	none	with coughing, sneezing, etc.	spontaneous, some control	no control (in the absence of fistula)	-
Operative injury to bladder and/or ureter	none	-	injury of bladder with primary repair	sepsis, fistula, or obstruction requiring secondary surgery; loss of one kidney; injury requiring anastomosis or re-implantation	septic obstruction of both kidneys or vesicovaginal fistula requiring diversion
Proteinuria	normal or < 0.15 g/24 hours	1+ or 0.15 - 1.0 g/24 hours	2+ to 3+ or 1.0 - 3.5 g/24 hours	4+ or > 3.5 g/24 hours	nephrotic syndrome
Note: If there is an inconsister	ncy between absolute v	alue and uristix reading, use	the absolute value for gradin	ıg.	
Renal failure	none	-	-	requiring dialysis, but reversible	requiring dialysis and irreversible
Ureteral obstruction	none	unilateral, not requiring surgery	-	bilateral, not requiring surgery	stent, nephrostomy tube, or surgery
Urinary electrolyte wasting (e.g., Fanconi's syndrome, renal tubular acidosis)	none	asymptomatic, not requiring treatment	mild, reversible and manageable with oral replacement	reversible but requiring IV replacement	irreversible, requiring continued replacement
Also consider Acidosis, Bicar	bonate, Hypocalcemia,	Hypophosphatemia.			
Urinary frequency/urgency	normal	increase in frequency or nocturia up to 2 x normal	increase > 2 x normal but < hourly	hourly or more with urgency, or requiring catheter	-
Urinary retention	normal	hesitancy or dribbling, but no significant residual urine; retention occurring during the immediate postoperative period	hesitancy requiring medication or occasional in/out catheterization (<4 x per week), or operative bladder atony requiring indwelling catheter beyond immediate postoperative period but for < 6 weeks	requiring frequent in/out catheterization (≥ 4 x per week) or urological intervention (e.g., TURP, suprapubic tube, urethrotomy)	bladder rupture

		G	rade		
Toxicity	0	1	2	3	4
Urine color change (not related to other dietary or physiologic cause e.g., bilirubin, concentrated urine, hematuria)	normal	asymptomatic, change in urine color	-	-	-
Vaginal bleeding is graded in t	the HEMORRHA	GE category.			
Vaginitis (not due to infection)	none	mild, not requiring treatment	moderate, relieved with treatment	severe, not relieved with treatment, or ulceration not requiring surgery	ulceration requiring surgery
Renal/Genitourinary-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling
		SECONDARY	MALIGNANCY		
Secondary Malignancy- Other (Specify type,) excludes metastastic tumors	none	-	-	-	present
		SEXUAL/REPROD	UCTIVE FUNCTI	ION	
Dyspareunia is graded in the P	AIN category.				
Dysmenorrhea is graded in the	PAIN category.				
Erectile impotence	normal	mild (erections impaired but satisfactory)	moderate (erections impaired, unsatisfactory for intercourse)	no erections	-
Female sterility	normal	-	-	sterile	-
Femininization of male is grad	led in the ENDOC	CRINE category.			
Irregular menses (change from baseline)	normal	occasionally irregular or lengthened interval, but continuing menstrual cycles	very irregular, but continuing menstrual cycles	persistent amenorrhea	-
Libido	normal	decrease in interest	severe loss of interest	-	-
Male infertility	-	-	Oligospermia (low sperm count)	Azoospermia (no sperm)	-
Masculinization of female is g	raded in the END	OCRINE category.			
Vaginal dryness	normal	mild	requiring treatment and/or interfering with sexual function, dyspareunia	-	-
Sexual/Reproductive Function-Other (Specify,)	none	mild	moderate	severe	disabling
	SYN	DROMES (not inclu	ded in previous ca	tegories)	
Acute vascular leak syndrome	is graded in the C	CARDIOVASCULAR (GENERA	L) category.		
	-	· · · · · · · · · · · · · · · · · · ·			

Grade							
Toxicity	0	1	2	3	4		
Autoimmune reactions are	e graded in the ALLE	ERGY/IMMUNOLOGY category.					
DIC (disseminated intrava	ascular coagulation) i	s graded in the COAGULATION c	ategory.				
Fanconi's syndrome is gra	aded as Urinary electr	rolyte wasting in the RENAL/GEN	ITOURINARY category.				
Renal tubular acidosis is g	graded as Urinary ele	ctrolyte wasting in the RENAL/GE	NITOURINARY category.				
Stevens-Johnson syndrom	ne (erythema multifor	me) is graded in the DERMATOLO	OGY/SKIN category.				
SIADH (syndrome of inap	ppropriate antidiuretion	c hormone) is graded in the ENDO	CRINE category.				
Thrombotic microangiopa	athy (e.g., thromboitic	c thrombocytopenic purpura/TTP or	r hemolytic uremic syndrom	n/HUS) is graded in the CO	AGULATION category.		
Tumor flare	none	mild pain not interfering with function	moderate pain; pain or analgesics interfering with function, but not interfering with activities of daily living	severe pain; pain or analgesics interfering with function and interfering with activities of daily living	Disabling		
Also consider Hypercalce	mia.						
Note: Tumor flare is characterized by a constellation of symptoms and signs in direct relation to initiation of therapy (e.g., anti-estrogens/androgens or additional hormones). The symptoms/signs include tumor pain, inflammation of visible tumor, hypercalcemia, diffuse bone pain, and other electrolyte disturbances.							
Tumor lysis syndrome	absent	-	-	present	-		
Also consider Hyperkalen	nia, Creatinine.						
Urinary electrolyte wastin	ng (e.g., Fanconi's syr	ndrome, renal tubular acidosis) is g	raded under the RENAL/GI	ENITOURINARY category.			
Syndromes-Other (Specify,)	none	mild	moderate	severe	life-threatening or disabling		

Appendix I

Toxicity Module

To be implemented at the request of the study sponsor or principal investigator in the protocol or by protocol amendment when more detailed information is considered pertinent.

Toxicity:	Date of Treatment:		Course Number:
Date of onset:			Grade at onset:
Date of first change in grade:			Grade:
Date of next change in grade:			Grade:
Date of next change in grade:			Grade:
Date of next change in grade:			Grade:
Date of next change in grade:			Grade:
Date of next change in grade:			Grade:
Did toxicity resolve? If so, date of resolution of toxicity:	Yes	No	
Date of last observation (if prior to recovery):			
Reason(s) observations stopped (if prior to recovery):			
Was patient retreated?	Yes	No	
If yes, was treatment delayed for recovery?	Yes	No	
Date of next treatment?		1,0	
Dose reduced for next treatment?	Yes	No	
Additional Comments:			
If module is being activated for new toxici	ty not currently in CTC, pla	assa provida dafir	nitions for toxicity grading:
Grade 0 =	-	_	initions for toxicity grading.
·			
Grade 1 = Grade 2 =			
Grade 3 =			
Grade 4 =			

Appendix II

Infection Module

To be implemented at the request of the study sponsor or principal investigator in the protocol or by protocol amendment when more detailed information is considered pertinent.

1.	. Use the Common Toxicity Criteria definitions to grade the severity of the infection.								
2.	Specify type of in	fection from the	following (CHOOSE O	NE):					
	BACTERIAL	FUNGAL	PROTOZOAL	VIRAL	UNKNOWN				
3.	Specify site of infe	ection from the f	ollowing (CHOOSE AL	L THAT APPLY	·):				
	BLOOD CULTUI BONE INFECTIO CATHETER (intr CATHETER (intr CENTRAL NERV EAR INFECTION EYE INFECTION GASTROINTEST ORAL INFECTION PNEUMONIA SKIN INFECTION UPPER RESPIRA URINARY TRAC VAGINAL INFECION, not	ON avenous) avenous), tunnel VOUS SYSTEM I I II INAL INFECTION I I I I I I I I I I I I I I I I I I	INFECTION ON)					
4.	Specify organism,	if known:	·						
5.	Prophylactic antib	iotic, antifungal,	or antiviral therapy adm	ninistration					
	Yes	No							
	If prophylaxis was	s given prior to in	nfection, please specify	below:					
	Antibiotic prophy	laxis							
	Antifungal prophy	laxis							
	Antiviral prophyla	axis							
	Other prophylaxis								

Appendix III

Performance Status Scales/Scores

ECOG or Zubrod scale		Karnofsky score
0	Asymptomatic and fully active	100%
1	Symptomatic; fully ambulatory; restricted in physically strenuous activity	80-90%
2	Symptomatic; ambulatory; capable of self-care; more than 50% of waking hours are spent out of bed	60-70%
3	Symptomatic; limited self-care; spends more than 50% of time in bed, but not bedridden	40-50%
4	Completely disabled; no self-care; bedridden	20-30%

Appendix IV

RTOG/EORTC Late Radiation Morbidity Scoring Scheme

Use for toxicities occuring greater than 90 days after radiation therapy.

Grade							
Toxicity	0	1	2	3	4		
Bladder- Late RT Morbidity Scoring	No change from baseline	Slight epithelial atrophy/minor telangiectasia (microscopic hematuria)	Moderate frequency/ generalized telangiectasia/ intermittent macroscopic hematuria	Severe frequency and dysuria/severe generalized telangiectasia (often with petechiae); frequent hematuria; reduction in bladder capacity (< 150 cc)	Necrosis/contracted bladder (capacity < 100 cc)/severe hemorrhagic cystitis		
Bone- Late RT Morbidity Scoring	No change from baseline	Asymptomatic; no growth retardation; reduced bone density	Moderate pain or tenderness; growth retardation; irregular bone sclerosis	Severe pain or tenderness; complete arrest of bone growth; dense bone sclerosis	Necrosis/ spontaneous fracture		
Brain- Late RT Morbidity Scoring	No change from baseline	Mild headache; slight lethargy	Moderate headache; great lethargy	Severe headaches; severe CNS dysfunction (partial loss of power or dyskinesia)	Seizures or paralysis; coma		
Esophagus- Late RT Morbidity Scoring	No change from baseline	Mild fibrosis; slight difficulty in swallowing solids; no pain on swallowing	Unable to take solid food normally; swallowing semi-solid food; dilatation may be indicated	Severe fibrosis; able to swallow only liquids; may have pain on swallowing; dilation required	Necrosis/ perforation; fistula		
Heart- Late RT Morbidity Scoring	No change from baseline	Asymptomatic or mild symptoms; transient T wave inversion and ST changes; sinus tachycardia > 110 (at rest)	Moderate angina on effort; mild pericarditis; normal heart size; persistent abnormal T wave and ST changes; low QRS	Severe angina; pericardial effusion; constrictive pericarditis; moderate heart failure; cardiac enlargement; EKG abnormalities	Tamponade/severe heart failure/severe constrictive pericarditis		
Joint- Late RT Morbidity Scoring	No change from baseline	Mild joint stiffness; slight limitation of movement	Moderate stiffness; intermittent or moderate joint pain; moderate limitation of movement	Severe joint stiffness; pain with severe limitation of movement	Necrosis/complete fixation		
Kidney- Late RT Morbidity Scoring	No change from baseline	Transient albuminuria; no hypertension; mild impairment of renal function; urea 25 - 35 mg%; creatinine 1.5 - 2.0 mg%; creatinine clearance > 75%	Persistent moderate albuminuria (2+); mild hypertension; no related anemia; moderate impairment of renal function; urea > 36 - 60 mg%; creatinine clearance > 50 - 74%	Severe albuminuria; severe hypertension; persistent anemia (< 10 g%); severe renal failure; urea > 60 mg%; creatinine > 4 mg%; creatinine clearance < 50%	Malignant hypertension; uremic coma/urea > 100%		
Larynx- Late RT Morbidity Scoring	No change from baseline	Hoarseness; slight arytenoid edema	Moderate arytenoid edema; chondritis	Severe edema; severe chondritis	Necrosis		

Appendix IV (continued)

RTOG/EORTC Late Radiation Morbidity Scoring Scheme

Use for toxicities occuring greater than 90 days after radiation therapy.

	Grade							
Toxicity	0	1	2	3	4			
Liver- Late RT Morbidity Scoring	No change from baseline	Mild lassitude; nausea; dyspepsia; slightly abnormal liver function	Moderate symptoms; some abnormal liver function tests; serum albumin normal	Disabling hepatic insufficiency; liver function tests grossly abnormal; low albumin; edema or ascites	Necrosis/hepatic coma or encephalopathy			
Lung- Late RT Morbidity Scoring	No change from baseline	Asymptomatic or mild symptoms (dry cough); slight radiographic appearances	Moderate symptomatic fibrosis or pneumonitis (severe cough); low grade fever; patchy radiographic appearances	Severe symptomatic fibrosis or pneumonitis; dense radiographic changes	Severe respiratory insufficiency/ continuous O ₂ /assisted ventilation			
Mucous membrane- Late RT Morbidity Scoring	No change from baseline	Slight atrophy and dryness	Moderate atrophy and telangiectasia; little mucus	Marked atrophy with complete dryness; severe telangiectasia	Ulceration			
Salivary glands- Late RT Morbidity Scoring	No change from baseline	Slight dryness of mouth; good response on stimulation	Moderate dryness of mouth; poor response on stimulation	Complete dryness of mouth; no response on stimulation	Fibrosis			
Skin- Late RT Morbidity Scoring	No change from baseline	Slight atrophy; pigmentation change; some hair loss	Patchy atrophy; moderate telangiectasia; total hair loss	Marked atrophy; gross telangiectasia	Ulceration			
Small/Large intestine- Late RT Morbidity Scoring	No change from baseline	Mild diarrhea; mild cramping; bowel movement 5 x daily slight rectal discharge or bleeding	Moderate diarrhea and colic; bowel movement > 5 x daily; excessive rectal mucus or intermittent bleeding	Obstruction or bleeding, requiring surgery	Necrosis/ perforation fistula			
Spinal cord- Late RT Morbidity Scoring	No change from baseline	Mild Lhermitte's syndrome	Severe Lhermitte's syndrome	Objective neurological findings at or below cord level treatment	Mono-, para-, quadriplegia			
Subcutaneous tissue- Late RT Morbidity Scoring	No change from baseline	Slight induration (fibrosis) and loss of subcutaneous fat	Moderate fibrosis but asymptomatic; slight field contracture; < 10% linear reduction	Severe induration and loss of subcutaneous tissue; field contracture > 10% linear measurement	Necrosis			
Eye- Late RT Morbidity Scoring	No change from baseline	Asymptomatic cataract; minor corneal ulceration or keratitis	Symptomatic cataract; moderate corneal ulceration; minor retinopathy or glaucoma	Severe keratitis; severe retinopathy or detachment; severe glaucoma	Panophthalmitis; blindness			
Radiation-Other (Specify,)	None	Mild	Moderate	Severe	Life-threatening or disabling			

Appendix V

BMT Complex/Multi-Component Events

Grade								
Toxicity	0	1	2	3	4			
	Note: The grading of Complex/Multi-Component Events in bone marrow transplant will be defined in the protocol. The grading scale must use the CTC criteria for grading the specific component events (toxicities).							
Failure to engraft	absent	mild	moderate	severe	life-threatening			
Also consider Hemoglobin (F	Igb), Neutrophils/granulo	ocytes (ANC/AGC), Platele	ts					
Graft versus host disease	absent	mild	moderate	severe	life-threatening			
Also consider Fatigue, Rash/o	desquamation, Diarrhea,	Bilirubin-GVHD						
Stem cell infusion complications	absent	mild	moderate	severe	life-threatening			
Also consider Allergic reaction/hypersensitivity, Arrhythmia, Hypertension, Hypotension, Fever, Rigors/chills, Sweating, Rash/desquamation, Urticaria, Diarrhea, Nausea, Vomiting, Hemorrhage/bleeding with grade 3 or 4 thrombocytopenia, Hemorrhage/bleeding without grade 3 or 4 thrombocytopenia, Hemoptysis, Alkaline phosphatase, Bilirubin, GGT, SGOT, SGPT, Infection, Hyperkalemia, Hypernatremia, Hypokalemia, Depressed level of consciousness, Seizures, Abdominal pain, Headache, Creatinine, Hemoglobinuria								
Veno-Occlusive Disease (VOD)	absent	mild	moderate	severe	life-threatening			
Also consider Weight gain-VOD, Bilirubin, Depressed level of consciousness, Hepatic pain, Renal failure, Hepatic enlargement.								